Aneurysmal Bone Cyst of the Paranasal Sinuses: A Rare Entity

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ABSTRACT

Aneurysmal bone cysts are vascular tumors that cause expansion and erosion of bone. We present this rare entity involving maxillary, ethmoid and frontal sinuses along with orbit. Computed tomography and histopathology showed typical features of aneurysmal bone cyst. We discuss clinical, radiological and histological details along with endoscopic management and literature review. The rarity of tumor makes this case interesting for head and neck surgeons.

Keywords: Aneurysmal bone cyst, Paranasal sinuses, Endoscopic management.

INTRODUCTION

Aneurysmal bone cysts (ABC) typically involve the long bones of the extremities, membranous bones of the thorax and pelvis, or vertebrae. The midline of the skull base is not the site of predilection for ABC and involvement of sinuses is extremely rare. There have been only five cases reports of involvement of sphenoid, ethmoid and maxillary sinuses.1-5 Skull and facial bone involvement accounts for 3 to 6% cases.3,4 The mandible is the most common site in head and neck region. Aneurysmal bone cyst may be found with other benign bone lesions, such as nonossifying fibroma, giant cell granuloma, fibrous dysplasia and fibromyxoma. Although ABC is non-neoplastic condition, its potential for rapid growth, considerable destruction of bone, and extension into adjacent structures warrants early intervention.6 The rarity of this disease in head and neck region and involvement of adjacent structures makes this case more interesting for head and neck surgeons.

CASE REPORT

A 12-year-old boy came to the ENT services of Outpatient Department of Postgraduate Institute of Medical Education and Research with complaints of right sided nasal obstruction for 7 months, pain and bulging of right eye for 6 months. Nasal obstruction was gradually progressing, initially on right nostril and later on left side. History of mouth breathing was present. There was proptosis associated with intermittent nonpurulent watery discharge from both eyes. Vision was normal. There was no history of trauma or nasal bleeding.

The general and systemic examinations were normal. On anterior rhinoscopy, a reddish fleshy mass was seen in the right nasal cavity completely filling it. The surface of the mass was smooth, pinkish and it was firm but compressible. Sensation of touch was intact. The right nasal chamber revealed severe deviated nasal septum. Posterior rhinoscopy showed smooth bulge in nasophyrnx. There was obvious axial proptosis on right, the ocular movements were normal, and pupillary reaction and corneal sensation were intact. Fundus examination was normal. Routine blood, serum electrolytes, chest X-ray and urine analysis were normal.

Skiagram of paranasal sinuses showed a lesion of mixed radiodensity in the area of ethmoid and left maxillary antrum. Computed tomography of nose, orbit and sinuses showed that there was a well-defined expansile heterogeneous bony mass on right side of nasal cavity including anterior and posterior ethmoid, nasopharynx, maxillary antrum, frontal sinus. There was medial expansile shift of lateral nasal wall with proptosis of eye and nasal septum displaced to left side (Figs 1 and 2). Based on clinical and radiological findings, a provisional diagnosis of fibro-osseous lesion was made.

The patient was operated upon under general anesthesia through endoscopic approach. There was a bony hard mass...
filling right nasal cavity, pushing middle turbinate, lamina papyracea and nasal septum laterally up to choana, ethmoid, maxillary and floor of frontal sinus. Entire mass which was having bony hard shell and cystic areas in between was removed. Right side medicated nasal packing was done. The specimen removed was sent for histopathological examination. The patient had an uneventful postsurgical course. The pack was removed on 3rd postoperative day and patient discharged on 4th postoperative day.

Biopsy showed intact respiratory epithelium lining. There were cystic spaces devoid of any lining; cyst wall showed fibrosis and few multinucleate giant cells in surrounding tissue. The cyst cavity showed RBC at places. The adjacent areas showed fibrous lesion, which were ossifying at places, suggesting healing areas of aneurysmal bone cyst.

The histopathological examination revealed overall features of aneurysmal bone cyst (Fig. 3). The patient is being regularly followed up for last one year, postoperative CT and nasal endoscopic examination did not show any features of residual disease (Fig. 4).

DISCUSSION

ABCs are rare, vascular benign tumor-like lesions of bone, of unknown origin. They represent 1 to 2% of all primary tumors of the bone and occur most commonly in females under 20 years of age. In 70% of cases, it is located in metaphysis of long bones, only 2% are reported in head

Fig. 1: CT scan axial view showing right maxillary sinus expansion

Fig. 2: Coronal section of CT scan showing mass in nasal cavity extending to maxilla, ethmoid and frontal sinuses with orbital involvement on right side

Fig. 3: Photomicrograph showing cystic spaces, RBCs and adjacent fibrosed areas ossification at places

Fig. 4: CT scan (two weeks after surgery)
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and neck. Kaffe et al (1999) reported no difference in sex distribution, ratio of involvement of maxilla and mandible is 1:2.4, and 92% of lesions were in posterior part of jaw. ABC of PNS typically present with pressure symptoms, such as headache, nasal congestion, cranial nerve palsies, loss of vision and proptosis. The current view in the literature is that cranial lesions constitute a separate entity, classified as primary or secondary depending on the presence or absence of an associated condition (giant cell tumors, chondroblastoma, fibrous dysplasia, etc.). Hady et al reported a case of aneurysmal bone cyst and claimed it to be the first reported case of aneurysmal bone cyst of maxilla. Radiological appearances can be diagnostic. The lesions are radiolucent in 87%, radiopaque in 2% and of mixed opacity in 11%. Plain skull radiographs are characterized by either a single or multiple blown-out lytic lesions. CT usually reveals an expansion and thinning of overlying bone that may be multiiloculated and might be confused with mucocele, but peripheral enhancement with iodide eliminates this possibility. MRI typically shows multiple fluid levels characteristic of ABC, although not diagnostic (rarely occurs in giant cell tumors, fibrous dysplasia, simple bone cyst, chondroblastoma, telangiectatic osteosarcoma). Microscopic examination reveals blood-filled channels bordered by a thin layer of spindle-shaped endothelial cells surrounded by connective tissue containing numerous multinucleated giant cells. Hemosiderin laden macrophages and new bone formation are found within the stromal matrix. ABC most likely represents a degenerative process associated with other primary bone lesions. The presence of multiple giant cells may cause difficulty in differentiating the lesion histologically from osteoclastoma, fibrous dysplasia, ossifying hematomata and cavernous hemangioma of bone. The principal diagnostic error occurs if the histopathologist fails to appreciate the lining of the blood-filled spaces.

The ABC is a benign lesion of obscure pathogenesis, it grows rapidly with considerable destruction of bone and extension into adjacent tissue. Accepted treatment options include surgical excision, curettage, bone grafting, cryotherapy and radiotherapy.

This is one of the rare ABC case reports, which was removed completely via endoscopic transnasal excision without any residual disease left behind.

REFERENCES